WEST Search History

Hide Items Restore Clear Cancel

DATE: Thursday, June 16, 2005

Hide?	Set Name	<u>Query</u>	Hit Count
	DB=PGPB; T	HES=ASSIGNEE; PLUR=YES; OP=ADJ	
	L9	17 or L8	9380
	L8	L6 and treating adj3 disease	7684
	L7	L6 and treating adj3 disorder	5437
	L6	phosphatase and human	24072
	DB = USPT, US	SOC,EPAB,JPAB,DWPI; THES=ASSIGNEE; PLUR=1	YES; OP=ADJ
	L5	12 or L4	3976
	L4	L1 and treating adj3 disorder	2794
	L3	L1 and treating adj3 disordere	0
	L2	L1 and treating disease	1707
	L1	phosphatase and human	26530

END OF SEARCH HISTORY

FILE 'HOME' ENTERED AT 08:45:50 ON 16 JUN 2005

=> FIL STNGUIDE => s phosphatase and human 50764 FILE MEDLINE T.1 25001 FILE CAPLUS L2 14869 FILE SCISEARCH 1.3 L44237 FILE LIFESCI 42900 FILE BIOSIS L5 35604 FILE EMBASE L6

TOTAL FOR ALL FILES

173375 PHOSPHATASE AND HUMAN

=> s 17 and treat? and (disease or disorder) TOTAL FOR ALL FILES 14175 L7 AND TREAT? AND (DISEASE OR DISORDER)

=> s 114 not 2004-2005/py

TOTAL FOR ALL FILES

12200 L14 NOT 2004-2005/PY L21

=> dup rem 121

7538 DUP REM L21 (4662 DUPLICATES REMOVED) L22

=> d 1-10 ibib abs

L22 ANSWER 1 OF 7538 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation

ACCESSION NUMBER: 2004:198859 SCISEARCH Full-text

THE GENUINE ARTICLE: 773VF

TITLE: Pimecrolimus: A review Gupta A K (Reprint); Chow M AUTHOR:

CORPORATE SOURCE: Suite 6, 490 Wonderland Rd S, London, ON N6K 1L6, Canada

(Reprint); Sunnybrook & Womens Coll, Hlth Sci Ctr, Div Dermatol, Toronto, ON, Canada; Univ Toronto, Toronto, ON,

Canada; Mediprobe Labs Inc, Toronto, ON, Canada

COUNTRY OF AUTHOR: Canada

SOURCE: JOURNAL OF THE EUROPEAN ACADEMY OF DERMATOLOGY AND

VENEREOLOGY, (SEP 2006) Vol. 17, No. 5, pp. 493-503. Publisher: BLACKWELL PUBLISHING LTD, 9600 GARSINGTON RD,

OXFORD OX4 2DG, OXON, ENGLAND.

ISSN: 0926-9959.

DOCUMENT TYPE: General Review; Journal

LANGUAGE: English REFERENCE COUNT: 43

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

ÀΒ Pimecrolimus (SDZ ASM 981), an ascomycin derivative, is one of the new classes of immunomodulating macrolactarns and was specifically developed for the treatment of inflammatory skin diseases. The interest in pimecrolimus has been substantial because of its significant anti-inflammatory activity and immunomodulatory capabilities and its low systemic immunosuppressive potential The mechanism of action of pimecrolimus is the blockage of T cell activation. Pimecrolimus (like all ascomycins) is an immunophilin ligand, which binds specifically to the cytosolic receptor, immunophilin macrophilin-12. This pimecrolimus- macrophilin complex effectively inhibits the protein phosphatase calcineurin, by preventing calcineurin from dephosphorylating the nuclear factor of activated T cells (NF-AT), a transcription factor. This results in the blockage of signal transduction pathways in T cells and the inhibition of the synthesis of inflammatory cytokines, specifically Th1- and Th2-type cytokines. Pimecrolimus has also been shown to prevent the release of cytokines and proinflammatory mediators from mast cells. Several studies have evaluated the effectiveness of pimecrolimus as a treatment for skin diseases. In animal models of allergic contact dermatitis, topical pimecrolimus was found to be effective. In human studies of allergic contact dermatitis pimecrolimus demonstrated significantly more efficacy than the control treatment. As well, the effectiveness of pimecrolimus 0.6% cream was comparable to 0.1% betamethasone-17-valerate; however, pimecrolimus was not associated with any of the side effects characteristic of a topical steroid. Topical application of pimecrolimus is not associated with skin atrophy. Pimecrolimus is effective and safe in both children and adults with atopic dermatitis. When

pimecrolimus 1% cream has been applied to adult atopics, improvement has been observed as early as the first week, with a 72% reduction in severity after 3 weeks. Pharmacokinetic studies have shown very low blood levels of pimecrolimus following topical application, with no accumulation after repeated applications. Following application of pimecrolimus cream occasional transient irritation may be experienced at the application site. Similar results have also been found in children aged 3 months and older following application of pimecrolimus 1% cream. Topical pimecrolimus in psoriasis appears to exhibit a dose-dependent therapeutic effect under semiocclusive conditions. Pimecrolimus has an enormous potential as a new treatment of inflammatory skin diseases. It has been shown to be effective in atopic and allergic contact dermatitis, with a favorable adverse-effects profile, which includes little effect on the systemic immune response.

L22 ANSWER 2 OF 7538 MEDLINE on STN

ACCESSION NUMBER: 2003082608 MEDLINE Full-text

DOCUMENT NUMBER: PubMed ID: 12594123

TITLE: Zoledronate treatment in active Paget's

disease.

AUTHOR: Chung G: Keen R W

Annals of the rheumatic diseases, (2003 Mar) 62 (3) 275-6. SOURCE:

Journal code: 0372355. ISSN: 0003-4967.

PUB. COUNTRY: England: United Kingdom

DOCUMENT TYPE: (CASE REPORTS)

Letter

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200304

Entered STN: 20030221 ENTRY DATE:

> Last Updated on STN: 20030410 Entered Medline: 20030409

L22 ANSWER 3 OF 7538 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on

STN

ACCESSION NUMBER: 2004:26913 BIOSIS Full-text

DOCUMENT NUMBER: PREV200400028085

N-acetylcysteine in the treatment of TITLE:

non-alcoholic steatohepatitis.

AUTHOR (S): Pamuk, Gulsum Emel [Reprint Author]; Sonsuz, Abdullah

[Reprint Author]

Division of Hepatology, Department of Internal Medicine, CORPORATE SOURCE:

Cerrahpapa Medical Faculty, University of Istanbul,

Istanbul, Turkey

SOURCE: Journal of Gastroenterology and Hepatology, (October 2003)

Vol. 18, No. 10, pp. 1220-1221. print.

CODEN: JGHEEO. ISSN: 0815-9319.

DOCUMENT TYPE: Letter LANGUAGE: English

ENTRY DATE: Entered STN: 31 Dec 2003

Last Updated on STN: 31 Dec 2003

L22 ANSWER 4 OF 7538 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on

STN

ACCESSION NUMBER: 2003:409031 BIOSIS Full-text

DOCUMENT NUMBER: PREV200300409031

TITLE: Response: Abnormal ALP isoenzyme in children with epilepsy

treated with carbamazepine.

AUTHOR (S): Verrotti, Alberto [Reprint Author]; Greco, Rita [Reprint

Author]; Latini, Giuseppe; Morgese, Guido; Chiarelli,

Francesco [Reprint Author]

CORPORATE SOURCE: Department of Pediatrics, University of Chieti, Chieti,

Italy

SOURCE: Epilepsia, (August 2003) Vol. 44, No. 8, pp. 1129. print.

ISSN: 0013-9580 (ISSN print).

DOCUMENT TYPE: Letter LANGUAGE:

English

ENTRY DATE: Entered STN: 3 Sep 2003

Last Updated on STN: 3 Sep 2003

L22 ANSWER 5 OF 7538 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN

ACCESSION NUMBER: 2003:409030 BIOSIS Full-text

DOCUMENT NUMBER: PREV200300409030

TITLE: Abnormal ALP isoenzyme in children with epilepsy

treated with carbamazepine.

AUTHOR(S): Okazaki, Toshio [Reprint Author]; Suzuki, Mitsuyuki; Nagai,

Tatsuo (Reprint Author)

CORPORATE SOURCE: Department of Forensic Medicine and Science, Graduate

School of Medicine, Kitasato University, Kitasato,

Sagamihara-shi, Kanagawa Prefecture, Japan

SOURCE: Epilepsia, (August 2003) Vol. 44, No. 8, pp. 1128. print.

ISSN: 0013-9580 (ISSN print).

DOCUMENT TYPE:

PE: Letter English

LANGUAGE: English ENTRY DATE: Entered

Entered STN: 3 Sep 2003

Last Updated on STN: 3 Sep 2003

L22 ANSWER 6 OF 7538 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on

STN

ACCESSION NUMBER: 2003:421557 BIOSIS Full-text

DOCUMENT NUMBER: PREV200300421557

TITLE: Glipizide treatment with short-term alcohol abuse

resulting in subfulminant hepatic failure.

AUTHOR(S): Ilario, Marius John-Marc; Turyan, Hach Vladimir; Axiotis,

Constantine A. [Reprint Author]

CORPORATE SOURCE: Department of Pathology, Health Science Center at Brooklyn,

State University of New York, 450 Clarkson Avenue, Box 25,

Brooklyn, NY, 11203-2098, USA

axiotismd@aol.com

SOURCE: Virchows Archiv, (July 2003) Vol. 443, No. 1, pp. 104-105.

print.

ISSN: 0945-6317.

DOCUMENT TYPE:

Letter English

LANGUAGE: ENTRY DATE:

Entered STN: 10 Sep 2003

Last Updated on STN: 10 Sep 2003

L22 ANSWER 7 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

ACCESSION NUMBER:

2003:942767 CAPLUS Full-text

DOCUMENT NUMBER:

140:40262

TITLE: Genes expressed in atherosclerotic tissue and their

use in diagnosis and pharmacogenetics

INVENTOR(S):

Nevins, Joseph; West, Mike; Goldschmidt, Pascal Duke University, USA

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 408 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.										ICAT	DATE						
WO 2003091391			A2 20031106														
				AT, AU,					B, BG, BR,					_			
																	JP,
•				-		-	LC,					-	-		-		-
						-	PT,	-	-	-	-					-	-
				-	-	-	VN,	_	-	-	-	-	-	-	-	_	
		TJ,			-			-	-								-
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,	BE,	BG,
		CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,
		PT,	SE,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,
		NE,	SN,	TD,	TG												
WO	2003	0913	91		A2 20031106				WO 2002-US38221						20021112		
	W:	ΑE,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,
		DE,	DK,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IS,	JP,
		ΚE,	KG,	KP,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,	MN,
		MW,	MX,	NO,	ΝZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,
		TR,	TT,	UA,	UG,	UZ,	VN,	YU,	ZA,	ZW							
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	υG,	ZM,	ZW,	AM,	ΑŻ,	BY,
		KG,	KZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,

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FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                            US 2002-374547P
PRIORITY APPLN. INFO.:
                                                                P 20020423
                                                                P 20021024
                                            US 2002-420784P
                                            US 2002-421043P
                                                               P 20021025
                                                               P 20021108
A 20021112
                                            US 2002-424680P
                                            WO 2002-US38221
      Genes whose expression is correlated with an determinant of an atherosclerotic phenotype
ΔR
      are provided. Also provided are methods of using the subject atherosclerotic determinant
      genes in diagnosis and treatment methods, as well as drug screening methods. In addition,
       reagents and kits thereof that find use in practicing the subject methods are provided.
      Also provided are methods of determining whether a gene is correlated with a disease
      phenotype, where correlation is determined using a Bayesian anal. [This abstract record
       is one of three records for this document necessitated by the large number of index
       entries required to fully index the document and publication system constraints.).
L22 ANSWER 8 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
                        2004:2697 CAPLUS Full-text
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         140:53403
                         Methods of using benzothiophenone derivatives to
TITLE:
                         treat cancer or inflammation
INVENTOR(S):
                         Zhang, Zaihui; Daynard, Timothy S.; Kalmar, Gabriel
                         Bela
PATENT ASSIGNEE(S):
                         Kinetek Pharmaceuticals, Inc., Can.
                         PCT Int. Appl., 72 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO.
                         KIND DATE
                                            APPLICATION NO.
                                                                   DATE
                                             -----
                         A1
                                20031231
                                           WO 2003-CA921
     WO 2004000314
                                                                   20030618
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR,
             TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                            US 2002-390589P
                                                                P 20020621
OTHER SOURCE(S):
                         MARPAT 140:53403
      This invention is directed to the use of certain benzothiophenone derivs. in treating
AB
      hyperproliferative disorders, e.g., cancer, inflammation, etc. in a mammal. Of particular
       interest are hyperproliferative disorders associated with cellular modulation of protein
      phosphorylation states, i.e.. altered activity of phosphorylation modifying enzyme(s),
       e.g. protein kinases and protein phosphatases. In one aspect of the invention, compds.
       and pharmaceutical compns. of the invention are used to inhibit the activity of PTPN12 and
       PTPN2; these enzymes have been associated with alterations in the phosphorylation state of
       cellular proteins.
REFERENCE COUNT:
                         3
                               THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L22 ANSWER 9 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
                         2003:1006684 CAPLUS Full-text
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         140:53443
                         Methods and systems for management of Alzheimer's
TITLE:
                         disease
INVENTOR(S):
                         Shalev, Alon
PATENT ASSIGNEE(S):
                         Brainsgate Ltd., Israel
                         PCT Int. Appl., 139 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
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FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

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PATENT NO.
                                   DATE
                                                   APPLICATION NO.
                                                                              DATE
                            KIND
                                     -----
                             ----
                                                   -----
     WO 2003105658
                             A2
                                     20031224
                                                   WO 2003-IL508
                                                                              20030613
          W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
               CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
               PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT,
               \mathtt{TZ}, \mathtt{UA}, \mathtt{UG}, \mathtt{US}, \mathtt{UZ}, \mathtt{VC}, \mathtt{VN}, \mathtt{YU}, \mathtt{ZA}, \mathtt{ZM}, \mathtt{ZW}
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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               FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
               BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                                   US 2002-388931P
                                                                          P 20020614
                                                                          A 20021114
                                                   US 2002-294310
AB
```

A method is provided for the treatment of Alzheimer's disease (AD). The method includes stimulating a sphenopalatine ganglion (SPG) of a subject so that the concentration of a substance in a brain of the subject changes.

L22 ANSWER 10 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:855798 CAPLUS Full-text

DOCUMENT NUMBER:

139:333135

TITLE:

Combination therapy including a PPAR α/γ dual agonist, and use in the treatment of hyperglycemia, lipid disorders, and obesity in patients with type 2 diabetes or related

disorders

INVENTOR(S):

Moller, David E.; Wright, Samuel D.

PATENT ASSIGNEE(S): SOURCE:

Merck & Co., Inc., USA PCT Int. Appl., 37 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                         KIND
                               DATE
                                            APPLICATION NO.
                                                                    DATE
                                -----
                                            -----
     WO 2003088962
                         A1
                                20031030
                                            WO 2003-US11896
                                                                    20030415
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS,
            LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
        UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                            US 2002-373091P
                                                                P 20020416
                                                               P 20020607
                                            US 2002-387031P
```

The invention provides pharmaceutical compns. comprising a combination of a first drug AB which is a PPAR α/γ dual agonist and a second drug selected from (1) a cholesterol absorption inhibitor, (2) an HMG-CoA reductase inhibitor, (3) a bile acid sequestrant, (4) nicotinyl alc., nicotinic acid, or a salt thereof, (5) a PPARa agonist, (6) a phenolic antioxidant, (7) an acyl CoA-cholesterol acyltransferase (ACAT) inhibitor, and (8) a cholesterol ester transfer protein (CETP) inhibitor, including pharmaceutically acceptable salts of one or more of the active ingredients, and a pharmaceutically acceptable carrier. Such combinations are useful for treating hyperglycemia, lipid disorders , and obesity in patients who have type 2 diabetes, metabolic syndrome, insulin resistance, and impaired glucose tolerance.

REFERENCE COUNT:

THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 11-20 ibib abs

L22 ANSWER 11 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:737908 CAPLUS Full-text

DOCUMENT NUMBER:

139:256390

TITLE:

Transcriptional regulation of protein tyrosine phosphatase PTP-1B gene and its use for drug screening and therapy for diabetes and obesity

Fukada, Toshiyuki

INVENTOR(S): PATENT ASSIGNEE(S):

Cold Spring Harbor Laboratory, USA; Tonks, Nicholas K.

SOURCE:

PCT Int. Appl., 115 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE :

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO).	KIND	DATE	APPLICATION NO.	DATE				
		- 							
WO 200307	76634	A2	20030918	WO 2003-EP2552	20030312				
W: A	E, AG, AL,	AM, AT	, AU, AZ,	BA, BB, BG, BR, BY,	BZ, CA, CH, CN,				
C	CO, CR, CU,	CZ, DE	, DK, DM,	DZ, EC, EE, ES, FI,	GB, GD, GE, GH,				
G	M, HR, HU,	ID, IL	, IN, IS,	JP, KE, KG, KP, KR,	KZ, LC, LK, LR,				
I	s, LT, LU,	LV, MA	, MD, MG,	MK, MN, MW, MX, MZ,	NI, NO, NZ, OM,				
P	H, PL, PT,	RO, RU	, SC, SD,	SE, SG, SK, SL, TJ,	TM, TN, TR, TT,				
T	Z, UA, UG,	US, UZ	, VC, VN,	YU, ZA, ZM, ZW					
RW: G	H, GM, KE,	LS, MW	, MZ, SD,	SL, SZ, TZ, UG, ZM,	ZW, AM, AZ, BY,				
K	KG, KZ, MD,	RU, TJ	, TM, AT,	BE, BG, CH, CY, CZ,	DE, DK, EE, ES,				
F	I, FR, GB,	GR, HU	, IE, IT,	LU, MC, NL, PT, RO,	SE, SI, SK, TR,				
E	BF, BJ, CF,	CG, CI	, CM, GA,	GN, GQ, GW, ML, MR,	NE, SN, TD, TG				
US 200322	A1	20031204	US 2003-388215	20030311					
PRIORITY APPLN	I. INFO.:			US 2002-363787P	P 20020312				
				US 2002-435587P	P 20021220				
				US 2003-388215	A 20030311				

AB Compns. and methods relating to PTP1B associated disorders are provided, based on the discovery that a Y-box protein binding site is present as a transcription enhancer sequence in the promoter region situated upstream (i. e., 5' to) of the human PTPIB gene. This site, situated at nucleotides -155 through -132 of the human PTP1B gene, mediates specific binding interactions with the YB-1 transcription regulatory factor, a member of the Y-box family of proteins. YB-1-targeted antisense constructs reduced PTPlB expression levels, providing an alternative to PTP1B active-site directed regulation of PTP1B activity. Increased phosphorylation of insulin receptor was observed in Ratl cells that were transfected with YB-1 antisense RNA. The invention claims nucleic acid sequences for Y box protein binding site and for transcription factor YB-1. The invention also claims protein sequences for YB-1. The invention further claims methods for use of transcriptional regulation of human PTP-1 gene for screening drugs and for therapeutic treatment of diabetes and obesity. The invention includes use of reporter genes for measuring PTP-1B enhancer activity, antibodies to the Y-box protein, methods for identifying agents that impair binding of a Y-box protein to a PTP-1B gene promoter Y box, and methods for measuring cellular responses to insulin and leptin.

L22 ANSWER 12 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:678608 CAPLUS Full-text

DOCUMENT NUMBER:

139:219271

TITLE:

Modification of defensins and their use in modulating

immune response and antimicrobial activity

INVENTOR(S):

Moss, Joel; Hirayama, Toshiya; Wada, Akiharo; Levine,

Rodney L.; Paone, Gregorino

PATENT ASSIGNEE(S):

The Government of the United States of America as Represented by the Secretary of the Department of Health and Human Services, USA; Nagasaki University

PCT Int. Appl., 62 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

SOURCE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE				
WO 2003070176	A2	20030828	WO 2003-US4649	20030218				
WO 2003070176	A3	20031127						
W: AE, AG, AL,	AM, AT	, AU, AZ,	BA, BB, BG, BR, BY, BZ,	CA, CH, CN,				
CO, CR, CU,	CZ, DE	, DK, DM,	DZ, EC, EE, ES, FI, GB,	GD, GE, GH,				

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LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                              US 2002-358504P P 20020219
       This disclosure provides modified antimicrobial agents, for example modified defensin
       polypeptides. In one embodiment, compns. including a modified arginine residue, such as
       an ADP-ribosylated and/or ribosylated alpha defensin polypeptide, are provided. Also
       provided are methods of modulating an immune response using the modified defensin
       polypeptides. In one embodiment, a method is provided for modulating an antimicrobial
       activity. In another embodiment, a method if provided for inhibiting a cytotoxic activity. Also disclosed are methods for treating diseases in a subject that are
       associated with an immune response, such as inflammatory and pulmonary diseases, using the
       disclosed modified defensin polypeptides.
L22 ANSWER 13 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                          2003:634041 CAPLUS Full-text
DOCUMENT NUMBER:
                          139:173802
TITLE:
                          KDR-associated phosphatase as target for
                          screening of angiogenesis modulators for therapeutic
                          use
INVENTOR(S):
                          Peters, Kevin Gene
PATENT ASSIGNEE(S):
                          The Procter & Gamble Company, USA
                          PCT Int. Appl., 80 pp.
SOURCE:
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                       DATE
                                 -----
                          ----
     WO 2003067220
                          A2 20030814 WO 2003-US4029
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                          CI, CM, GA, GN, GQ, GN, .....
Al 20030821 US 2002-327414 20021220
US 2002-355125P P 20020208
     US 2003158083
PRIORITY APPLN. INFO.:
       KDR-associated phosphatase is useful as a target to screen for agents useful for the
       treatment of angiogenesis mediated disorders.
L22 ANSWER 14 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                          2003:633897 CAPLUS Full-text
DOCUMENT NUMBER:
                          139:178697
                          Screening of human monoclonal antibodies
TITLE:
                          against cell surface coreceptor of HIV for diagnosis
```

Hua, Shaobing; Pauling, Michelle H.; Zhu, Li

APPLICATION NO.

and therapy

CODEN: PIXXD2

Patent English

KIND

Genetastix Corporation, USA

PCT Int. Appl., 150 pp.

DATE

INVENTOR(S):

DOCUMENT TYPE:

SOURCE:

PATENT ASSIGNEE(S):

FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO.

GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,

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20030814
                                            WO 2003-US3763
                                                                    20030207
     WO 2003066830
                          A2
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                20030904
                                            US 2002-71866
     US 2003165988
                          A1
                                                                   20020208
PRIORITY APPLN. INFO.:
                                             US 2002-71866
                                                                 A1 20020208
                                             US 2002-133978
                                                                A1 20020425
      Methods are provided for efficient, high throughput screening of antibody libraries
AB
       against protein targets, especially membrane proteins. In particular, methods are
       provided for screening a fully human antibody library against membrane proteins such as
       chemokine receptors in yeast. More particularly, a library of human single chain
       antibodies is screened against peptide fragments derived from extracellular domains of
       human CXCR4 and CCR5 resp. and high affinity monoclonal antibodies against CXCR4 and CCR5
       are selected. The antibodies can be used as prophylactics or therapeutics to prevent and
       treat HIV infection, cancer and other diseases or conditions, as well as for screening
       drugs and diagnosing diseases or conditions associated with interactions with membrane
L22 ANSWER 15 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                         2003:633497 CAPLUS Full-text
DOCUMENT NUMBER:
                         139:174286
TITLE:
                         Use of GLP-1 compound for treatment of
                         critically ill patients
INVENTOR(S):
                         Knudsen, Lotte Bjerre; Selmer, Johan; Hansen, Kristian
                         Tage
PATENT ASSIGNEE(S):
                         Novo Nordisk A/S, Den.
SOURCE:
                         PCT Int. Appl., 40 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                DATE
                                             APPLICATION NO.
                                                                    DATE
                         _ _ _ _
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     WO 2003066084
                         A1
                                20030814
                                            WO 2003-DK61
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     US 2003199445
                         A1 20031023
                                            US 2003-359324
                                                                    20030206
PRIORITY APPLN. INFO .:
                                            DK 2002-184
                                                                 A 20020207
                                             US 2002-359834P
                                                                 P 20020226
       Use of medicament for life saving treatment of critically ill patients SIRS patients, and
AB
       method of treatment. The medicament comprises a GLP-1 compound which effectively controls
       the blood glucose level.
REFERENCE COUNT:
                               THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L22 ANSWER 16 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
                         2003:590947 CAPLUS Full-text
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         139:148004
TITLE:
                         Digenic mutations of PPARy and PPP1R3A
                         associated with severe insulin resistance and type 2
                         diabetes and their use in the diagnosis and
```

treatment of diabetes

Barroso, Ines; Schafer, Alan J.; O'Rahilly, Stephen

INVENTOR(S):

O.; Waraham, Nicholas J.

PATENT ASSIGNEE(S): SOURCE:

Incyte Genomics, Inc., USA PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent LANGUAGE: English

PATENT INFORMATION:

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PATENT NO.
                                                                                                  KIND
                                                                                                                             DATE
                                                                                                                                                                              APPLICATION NO.
                                                                                                                                                                                                                                                                           DATE
                   WO 2003061583 A2
                                                                                                                              20030731 WO 2003-US1625
                                                                                                                                                                                                                                                                          20030117
                   W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO,
                                  CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR,
                                  HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
                                  LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU,
                                  SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,
                                  \mbox{VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, \mbox{TM} \label{eq:control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_control_contro
                   RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB,
                                  GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR
PRIORITY APPLN. INFO.:
                                                                                                                                                                          US 2002-PV350405
                                                                                                                                                                                                                                                                           20020118
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The invention provides mutated genes for peroxisome proliferator-activated receptor gamma (PPARY) and the glycogen-associated regulatory subunit of protein phosphatase-1 (PPPIR3A), PPARYFS and PPP1R3AFS, resp., and polynucleotides and proteins of PPARYFS and PPP1R3AFS that are expressed in diabetes. In particular, frameshift/premature stop mutations in two unlinked genes, PPARy and PPP1R3A, which are key regulators of lipid metabolism (adipocyte differentiation) and glycogen synthesis (muscle /liver mode of glycogen storage) resp., are found in type II diabetes patients with severe insulin resistance of two families. Specifically, the frameshift/premature stop mutation (A553AAAiT)fs.185(stop 186) in PPARY results in a mutation of K = lysine to M = methionine at amino acid position 185 of PPARy within the second zinc-finger (Zn) of the DNA-binding domain (DBD) together with a mutation of S = serine to stop codon (X) at position 186 of PPARy. And the frameshift/premature stop mutation (C1984AG)fs.662(stop 668) in PPP1R3A results in mutation of an N = asparagine to stop codon at amino acid position 668 of PPP1R3A. The family pedigree shows the concordance of features related to severe insulin resistance and type 2 diabetes and the presence of PPARy and PPP1R3A mutations. These mutations are confirmed at the protein level by related protein activity assays. Also disclosed are the summaries of clin. and biochem. characteristics of the frameshift mutation carriers for both PPARYFS and PPP1R3AFS in both family members. Thus, a model of interactions among genes that may underlie common human metabolic disorders such as type 2 diabetes is suggested. It also provides for the use of the DNA mutation, the protein, a polynucleotide encoding the protein, and antibodies that specifically bind the protein in various methods to diagnose, stage, treat, or monitor the treatment of diabetes.

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L22 ANSWER 17 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
                        2003:551645 CAPLUS Full-text
ACCESSION NUMBER:
```

DOCUMENT NUMBER:

139:80300 TITLE:

Sequences of a human phosphatidic acid

phosphatase 2C sequence homolog and uses in

diagnosis, therapy and drug screening

INVENTOR(S): Zhu. Zhimin

PATENT ASSIGNEE(S): Bayer Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 102 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO	KI	KIND DATE			APPLICATION NO.							DATE				
WO 200305	A	A1 20030717			WO 2003-EP55						20030107					
W: A	E, AG,	AL, AM	, AT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,		
C	O, CR,	CU, C2	, DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,		
G	M, HR,	HU, II	, IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	ΚZ,	LC,	LK,	LR,		
L	S, LT,	LU, LV	, MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,		
P	L, PT,	RO, RU	, sc,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	ΤZ,		
U	A, UG,	US, UZ	, vc,	VN,	YU,	ZA,	ZM,	ZW								
RW: G	H, GM,	KE, LS	, MW,	ΜZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	BY,		
K	G, KZ,	MD, RU	, TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,		

FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG P 20020107 PRIORITY APPLN. INFO.: US 2002-344798P

The invention provides protein and cDNA sequences of a novel human phosphatidic acid AB phosphatase 2C sequence homolog. The invention also provides reagents and methods of regulating a human phosphatidic acid phosphatase 2C sequence homolog. Reagents that regulate human phosphatidic acid phosphatase 2 and reagents which bind to human phosphatidic acid phosphatase 2 gene products can play a role in preventing, ameliorating,

or correcting dysfunctions or diseases including cancer, CNS disorders, and COPD. 10

REFERENCE COUNT:

THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 18 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:434595 CAPLUS Full-text

DOCUMENT NUMBER:

139:32331

TITLE:

Protein-protein interactions involved in signaling by

transforming growth factor-\$\beta\$ or TGF\$ family

members and uses thereof

INVENTOR(S):

Legrain, Pierre; Gauthier, Jean-Michel; Colland,

Frederic; Jacq, Xavier

PATENT ASSIGNEE (S):

Hybrigenics, Fr. SOURCE: PCT Int. Appl., 148 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE:

Patent English

PATENT INFORMATION:

PAT	PATENT NO.						KIND DATE			APPLICATION NO.						DATE				
WO :	2003045990 A2					20030605			WO 2002-EP13866					20021126						
W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,	co,			
	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,			
	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,			
	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,	PL,	PT,	RO,	RU,			
	SC,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,			
	VC,	VN,	YU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	TJ,	TM					
RW:	ΑT,	BE,	BF,	ВJ,	CF,	CG,	CH,	CI,	CM,	CY,	DE,	DK,	ES,	FI,	FR,	GΑ,	GB,			
	GR,	ΙE,	IT,	LU,	MC,	ML,	MR,	NE,	NL,	PT,	SE,	SN,	TD,	TG,	TR					
PRIORITY	RIORITY APPLN. INFO.:						US 2001-PV333348													
					US 2002-PV384537 20020531											531				
									US	3 20	02-P	V422	471		2	0021	030			

AB The present invention relates to protein-protein interactions involved in transforming growth factor-\(\beta\) (TGF\(\beta\)) disorders and/or diseases. More specifically, the present invention relates to complexes of polypeptides or polynucleotides encoding the polypeptides, fragments of the polypeptides, Selected Interacting Domains (SID®) which are identified due to the protein-protein interactions, methods for screening drugs for agents which modulate the interaction of proteins, and pharmaceutical compns. that are capable of modulating the protein-protein interactions. The invention claims polynucleotide and polypeptide sequences for SID® proteins. Some examples show effects of siRNA downregulation or overexpression of SID proteins on TGFB- and bone morphogenetic proteindependent reporter activity.

L22 ANSWER 19 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:417966 CAPLUS Full-text

DOCUMENT NUMBER:

139:5611

TITLE:

SOURCE:

Mkk3b protein for screening modulators of lymphocyte

activation and diagnosing/prognosing/treating

disorders associated with lymphocyte

dysfunction

INVENTOR (S):

Fu, Alan C.; Wu, Jun; Liao, Charlene X.; Mancebo,

PATENT ASSIGNEE(S):

Rigel Pharmaceuticals, Inc., USA

PCT Int. Appl., 94 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.

KIND DATE APPLICATION NO.

DATE

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WO 2003044529
                               20030530
                                            WO 2002-US36881
                                                                    20021118
                         A1
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, IE, IT, LU; MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                           US 2001-332441P
                                                                 P 20011116
      The present invention provides compns. and methods for modulating lymphocyte activation.
      Nucleic acids encoding proteins and proteins so encoded which are capable of modulating
      lymphocyte activation are provided. Compns. and methods for the treatment of disorders
       related to lymphocyte dysfunction or dysregulation are also provided. Prophylactics and
      methods for the prevention of such disorders are also provided. Also provided are compns.
      and methods for diagnostic and prognostic determination of such disorders. Further
      provided are assays for the identification of bioactive agents capable of modulating
       lymphocyte activation.
REFERENCE COUNT:
                               THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L22 ANSWER 20 OF 7538 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                         2003:335288 CAPLUS Full-text
DOCUMENT NUMBER:
                         138:349758
TITLE:
                         DNA sequence of promoter for human
                         sphingosine kinase 1 and uses
                         Kohama, Takafumi; Sugiura, Masako
INVENTOR(S):
PATENT ASSIGNEE (S):
                         Sankyo Company, Limited, Japan
SOURCE:
                         PCT Int. Appl., 35 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                DATE
                                            APPLICATION NO.
                                                                    DATE
     WO 2003035871
                                20030501
                                            WO 2002-JP10882
                         A1
                                                                    20021021
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     JP 2003199590
                                20030715
                                            JP 2002-307956
                          A2
PRIORITY APPLN. INFO.:
                                            JP 2001-325402
                                                                 A 20011023
      This invention provides DNA sequence of promoter for human sphingosine kinase 1. The
      expression level of reporter gene was enhanced when the expression was regulated under
       sphingosine kinase 1 promoter. The promoter provided in this invention can be used for
      diagnosis, treatment and screening the drugs for arteriosclerosis, diabetes, thrombosis,
      inflammation, immunopathy, allergy, cancer and cancer metastasis.
                               THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
```

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT